

FIG. 1

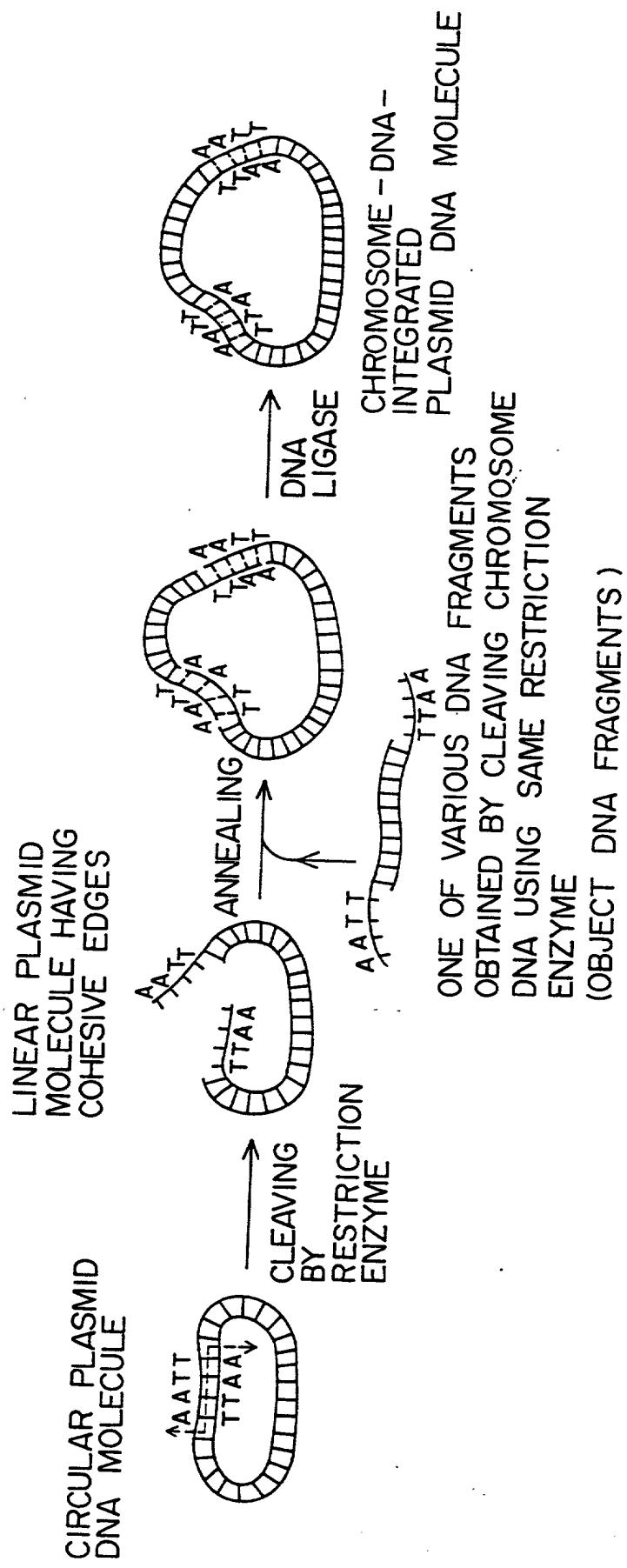


FIG. 2

SEQ ID: 1

SEQ ID: 2

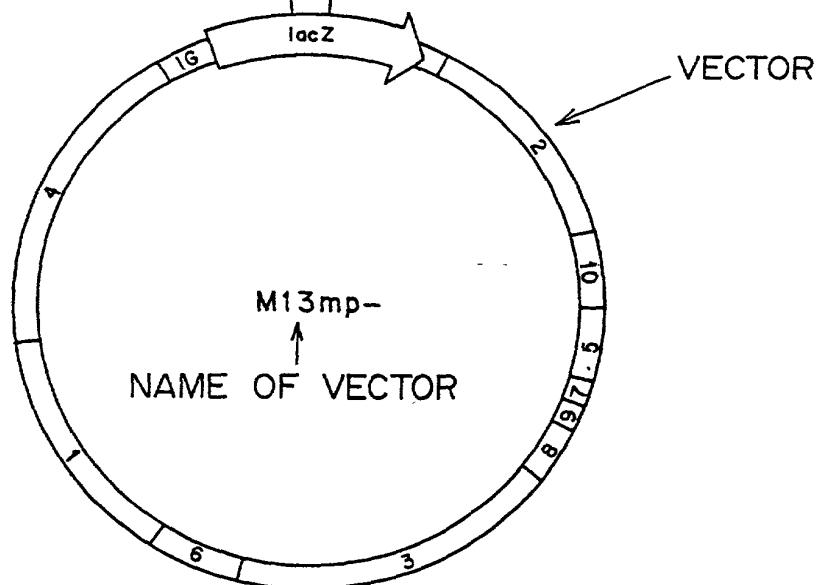
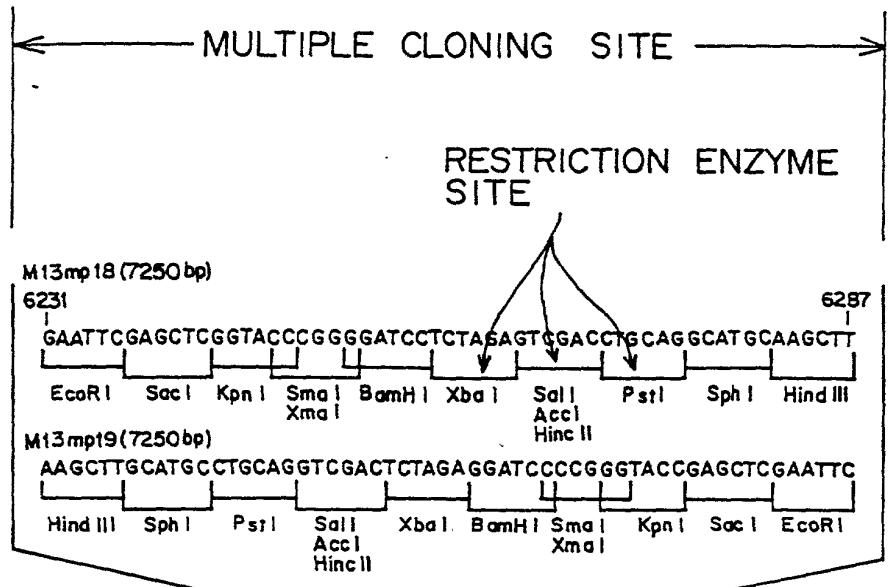
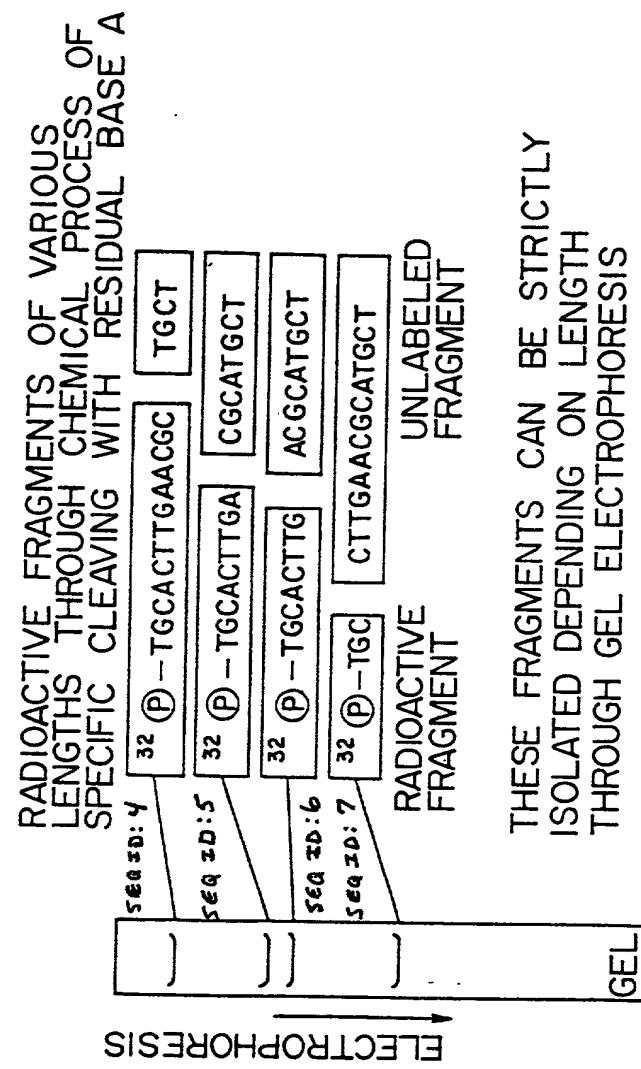


FIG. 3

DNA FRAGMENT LABELED WITH ^{32}P AT 5' EDGE
 $^{32}\text{P} - \text{TGCACCTAACGGCATGCT}$



THESE FRAGMENTS CAN BE STRICTLY ISOLATED DEPENDING ON LENGTH THROUGH GEL ELECTROPHORESIS

FIG. 4

RETRIEVAL KEY IS GENERATED TO RETRIEVE VECTOR UNIT
DEPENDING ON VECTOR AND RESTRICTION ENZYMES USED
ON VECTOR SIDE AND OBJECT DNA FRAGMENT SIDE

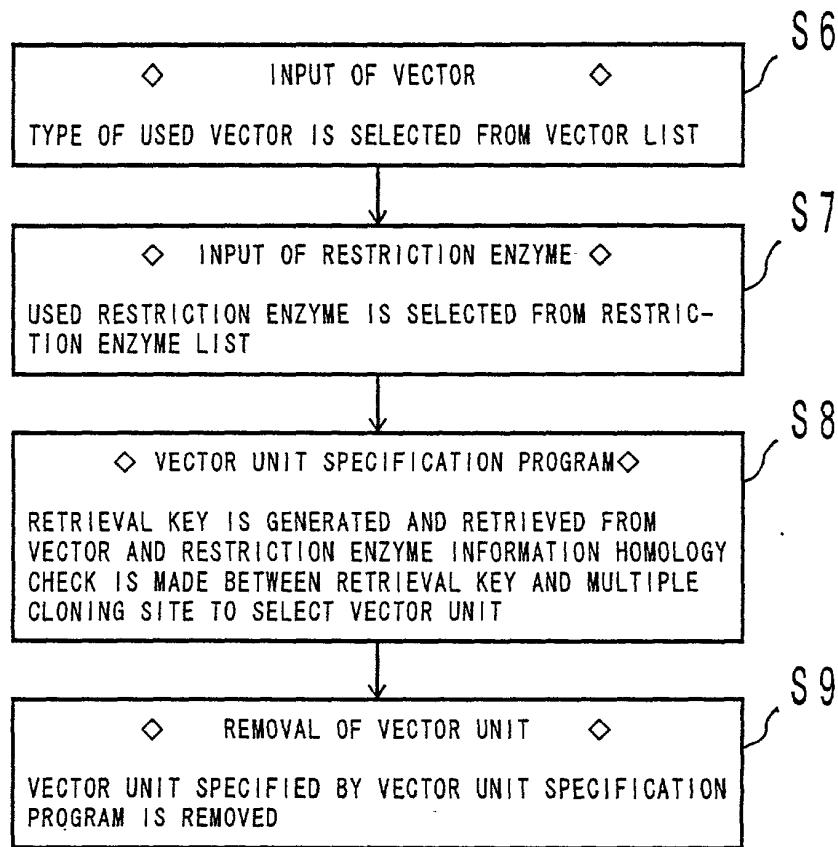
1

VECTOR UNIT IS SPECIFIED USING GENERATED RETRIEVAL
KEY AND AUTOMATICALLY REMOVED

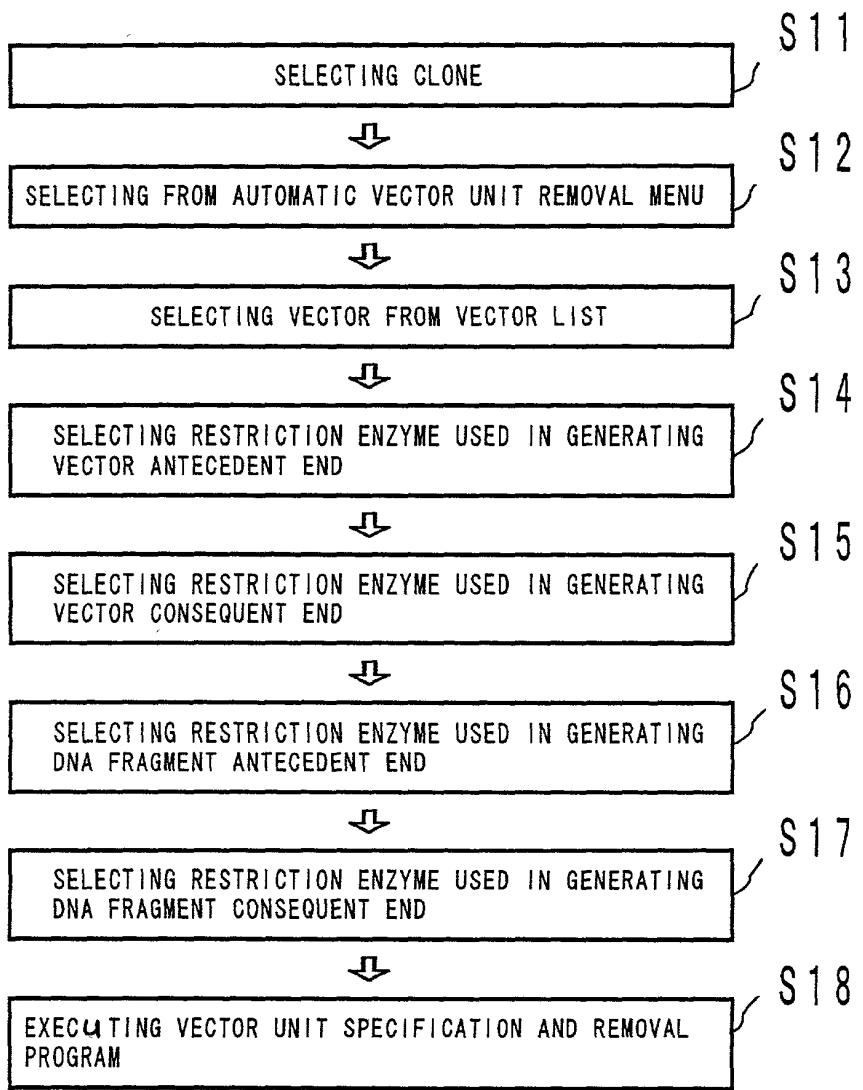
2

F I G . 5

9928640-0200-4



F I G. 6



F I G. 7

M13MP18
M13MP19
PBR322
PSL1180
PSL1190
PT7T318U
PT7T319U
PTZ18R
PTZ19R
PUC18
PUC19, ETC.

FIG. 8

VECTOR DB FORMAT

>ID
PUC18
>SEQ ID: 8

TCGCGCGTTGGTGTAGACGGTAAAAACCTCTGACACATGCAGCTCCGGAGACGGTCACAGCTTGTCTGTAAGCGGAT
GCCGGGAGCAGACAAGCCGTCAAGGGCGCGTCAGCGGGTGTGGCGGGTGTGGCTTAACATATGCGGCATCAGA
GCAGATTGACTGAGAGTCACCATATGCGGTGTGAAATACCGCACAGATGCGTAAGGAGAAAATACCGCATCAGCGCC
ATTCCGCAATTCAAGGCTGCGCACTGTGGGAAGGGCGATCGGTGCGGCCTCTCGCTATTACGCCAGCTGGCGAAAGGG
GGATGTGCTGCAAGGCAGTTAAGTGGGTACGCCAGGGTTTCCCAGTCACGACGTTGAAAACGACGGCCAGTGC
GCTTGATGCGCTGCAAGGTGACTCTAGAGGATCCCCGGTACCGAGCTGAATTGTAATCATGGTCAAGCTTTCT
GTGTGAAATTGTTATCCGCTACAATTCCACACACATACGAGCGGAAGCATAAAGTGTAAAGCTGGGTGCTTAATG
AGTGAGCTAACTCACATTAAATTGCGTTGCGCTACTGCCGCTTCCAGTCGGGAAACCTGTCGTGCCAGCTGCTTAAT
GAATCGGCCAACCGCGGGGAGAGGCGGGTTGCGTATTGGCGCTCTCCGCTTCCGCTACTGACTCGCTGCGCTCG
GTCGTGCGCTGCGCGAGCGGTATCAGCTACTCAAAGCGGTAAACGGTTATCCACAGAACTAGGGATAACGCA
AAAGAACATGTGAGCAAAAGGCCAGCAAAAGGCCAGGAACCGTAAAAGGCCGCGTTGCTGGCTTTCCATAGGCTCC
GCCCGCTGACGAGCATTACAAAAATGACGCTCAAGTCAGGGTGGCGAAACCCGACAGGACTATAAAAGATACCGCG
TTTCCCCCTGAAAGCTCCCTGCGCTCTCTGTTCCGACCCGCTTACCGGATACCTGTCGCCCTTCTCCCTC
GGGAAGCGTGGCGTTCTCAAAGCTACGCTGTAGGTATCTGAGTTGGTGTAGGTGCTCGCTTCAAGCTGGCTGTG
TGCACGAACCCCCCGTTAGCCGACCGCTGCCCTTACCGGTAACATATGCTCTGAGTCCAACCCGGTAAGACACGAC
TTATGCCACTGGCAGCAGCCACTGGTAACAGGATTAGCAGAGCAGGGTATGTAAGCGGTGCTACAGAGTTCTGAAGTG
GTGGCTAACTACGGCTACACTAGAAGAACAGTATTGGTATCTGCGCTCTGCTGAAGCCAGTTACCTCGGAAAAGAG
TTGGTAGCTTGTATCCGCAAAACAAACCACCGCTGGTAGCGGTGGTTTTTGTGTTGAAAGCAGCAGATTACGCG
AAAAAAGGATCTCAAGAACATCTTGTATCTTCTACGGGTCTGACGCTCAGTGGAACGAAAACACGTTAAGGGAT
TTGGTCATGAGATTATCAAAGGATCTCACCTAGATCCTTTAAATTAAAATGAAGTTTAAATCAATCTAAAGTA
TATATGAGTAAACTGGTCTGACAGTTACCAATGCTTAATCAGTGAGGACACCTATCTAGCAGTCTGCTATTGTTCA
TCCATAGTTGCCGACTCCCCGCTGTAGATAACTACGATACGGAGGGCTTACCATCTGGCCCCAGTGTGCAATGAT
ACCGCGAGACCCACGCTACCGCTCCAGATTATCAGCAATAACCAAGCCAGCGGAAGGGCGAGCGCAGAACGGTC
CTGCAACTTATCCGCTCCATCGCTATTAAATTGTCGGGAAGCTAGTAAGTAGTTGCCAGTTAATAGTTG
CGCAACGTTGTCATTGCTACAGGCATCGTGGTCACTGCTCGTGTGGTATGGCTTATTCACTCCGTTCCG
ACGATCAAGGCAGTTACATGATCCCCCATGGTGTGCAAAAAGCGGTAGCTCCTCGGCTCTCGATCGTTGTCAGAA
GTAAGTTGGCGCAGTGTATCACTCATGGTATGGCAGCACTGCATAATTCTTACTGTCATGCCATCCGTAAGATGC
TTTCTGTGACTGGTAGACTCAACCAAGTCATTCTGAGAAATAGTGTATGCGGCAGCGAGTTGCTCTGCCGGCGTC
AATACGGGATAATAACCGGCCACATAGCAGAACATTAAAGTGTCTCATCATTGGAAAACGTTCTCGGGCGAAAACCT
CAAGGATCTTACCGCTGGTAGAGATCCAGTCAGTGTAAACCCACTCGTGCACCCAACTGATCTCAGCATTTC
ACCAAGCGTTCTGGGTGAGCAAAAAGCGAAGGCAAAATGCCCAAAAAGGGAAATAAGGGCGACACGGAAATGTTGAAT
ACTCATACTCTTCTTTCAATATTATTGAAGCATTATCAGGGTTATTGCTCATGAGCGGATACATATTGAATGTA
TTAGAAAAATAACCAATAGGGGTTCCGCGCACATTCCCGAAAAGTGCACCTGACGTTAAGAAACCATTATTAC
ATGACATTAACCTATAAAATAGGCATACGAGGCCCTTCGTC

>MULTI
399..450

FIG. 9

(* INDICATES MULTIPLE CLONING SITE)

SEQ ID: 9 GTGCCAAGCTTGCATGCCCTGCAGGTCTGACTCTAGAGGATCCCCGGGTACCGAGCTCGAATTCTGTAAT

SEQ ID: 10 AAGCTT→HIND III

SEQ ID: 11 GCATGC→SPH I

SEQ ID: 12 CTGCAG→PST I

SEQ ID: 13 GTCGAC→SAL I, ACC I, HINC II

SEQ ID: 14 TCTAGA→XBA I

SEQ ID: 15 GGATCC→BAMH I

SEQ ID: 16 CCCGGG →SMA I, XMA I

SEQ ID: 17 GGTACC →KPN I

SEQ ID: 18 GAGCTC →SAC I

SEQ ID: 19 GAATTC →ECOR I

FIG. 10

VECTOR SIDE	OBJECT DNA FRAGMENT SIDE
HIND III	HINDIII
SPH I	SPH I
PST I	PST I
SAL I	SAL I
ACC I	ACC I
HINC II	HINC II
XBA I	XBA I
BAMH I	BAMH I
SMA I	SMA I
XMA I	XMA I
KPN I	KPN I
SAC I	SAC I
ECOR I	ECOR I
OTHER RESTRICTION ENZYME	
• • •	

FIG. 11

S 21

DETERMINING RETRIEVAL KEY

TWO RETRIEVAL KEYS ARE GENERATED ON EACH OF 5' (FORWARD) AND 3' (BACKWARD) SIDES ACCORDING TO VECTOR TYPE AND RESTRICTION ENZYME INFORMATION



S 22

HOMOLOGY RETRIEVAL

AFTER HOMOLOGY RETRIEVAL USING RETRIEVAL KEY,
PRIMARY CANDIDATE LISTS FOR BOUNDARY PORTION 5'
AND 3' SIDES ARE GENERATED



S 23

HOMOLOGY CHECK

HOMOLOGY CHECK IS MADE BETWEEN MULTIPLE CLONING
SITE AND PRECEDING AREA OF PRIMARY CANDIDATE FOR
5' BOUNDARY PORTION AND FOLLOWING AREA OF PRIMARY
CANDIDATE FOR 3' BOUNDARY PORTION TO GENERATE
LIST OF SECONDARY CANDIDATES FOR BOUNDARY PORTION



S 24

SPECIFYING BOUNDARY AREA

CHECK THAT EACH CANDIDATE IS UNIQUE, AND CHECK
POSITIONAL RELATIONSHIP BETWEEN 5' SIDE SECONDARY
CANDIDATE AND 3' SIDE SECONDARY CANDIDATE. IF OK,
THESE CANDIDATES ARE SPECIFIED AS VECTOR UNIT.



S 25

DETERMINING PORTION CLEAVED

PORTION CLEAVED IN BOUNDARY AREA IS DETERMINED

F I G. 1 2

WHEN SINGLE-STRANDED AREA IS FOUND ON 3' SIDE

STRAND A 5'	AREA A	AREA B3	AREA C	3'
STRAND B 3'	AREA C	AREA B3	AREA A	5'
	← RESTRICTION ENZYME →	SITE		

FIG. 13A

WHEN NO SINGLE-STRANDED AREA IS FOUND

STRAND A 5'	AREA A	AREA C	3'
STRAND B 3'	AREA C	AREA A	5'
	← RESTRICTION ENZYME →	SITE	

FIG. 13B

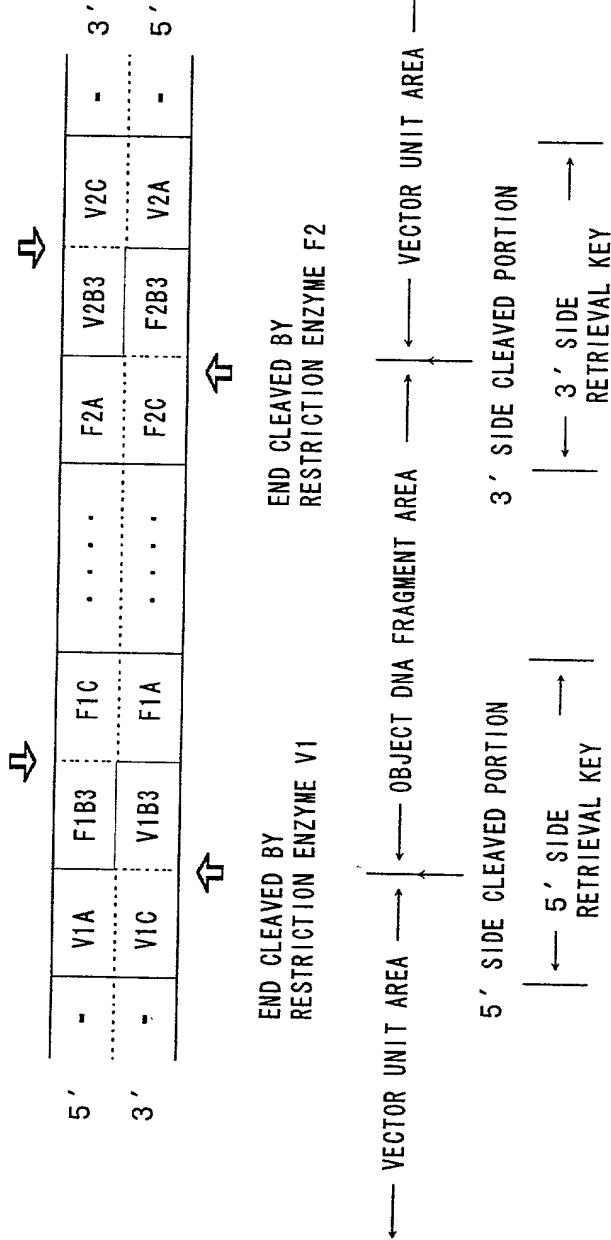
WHEN SINGLE-STRANDED AREA IS FOUND ON 5' SIDE

STRAND A 5'	AREA A	AREA B5	AREA C	3'
STRAND B 3'	AREA C	AREA B5	AREA A	5'
	← RESTRICTION ENZYME →	SITE		

FIG. 13C

END CLEAVED BY
RESTRICTION ENZYME F1

END CLEAVED BY
RESTRICTION ENZYME V2



F 1 G. 1 4

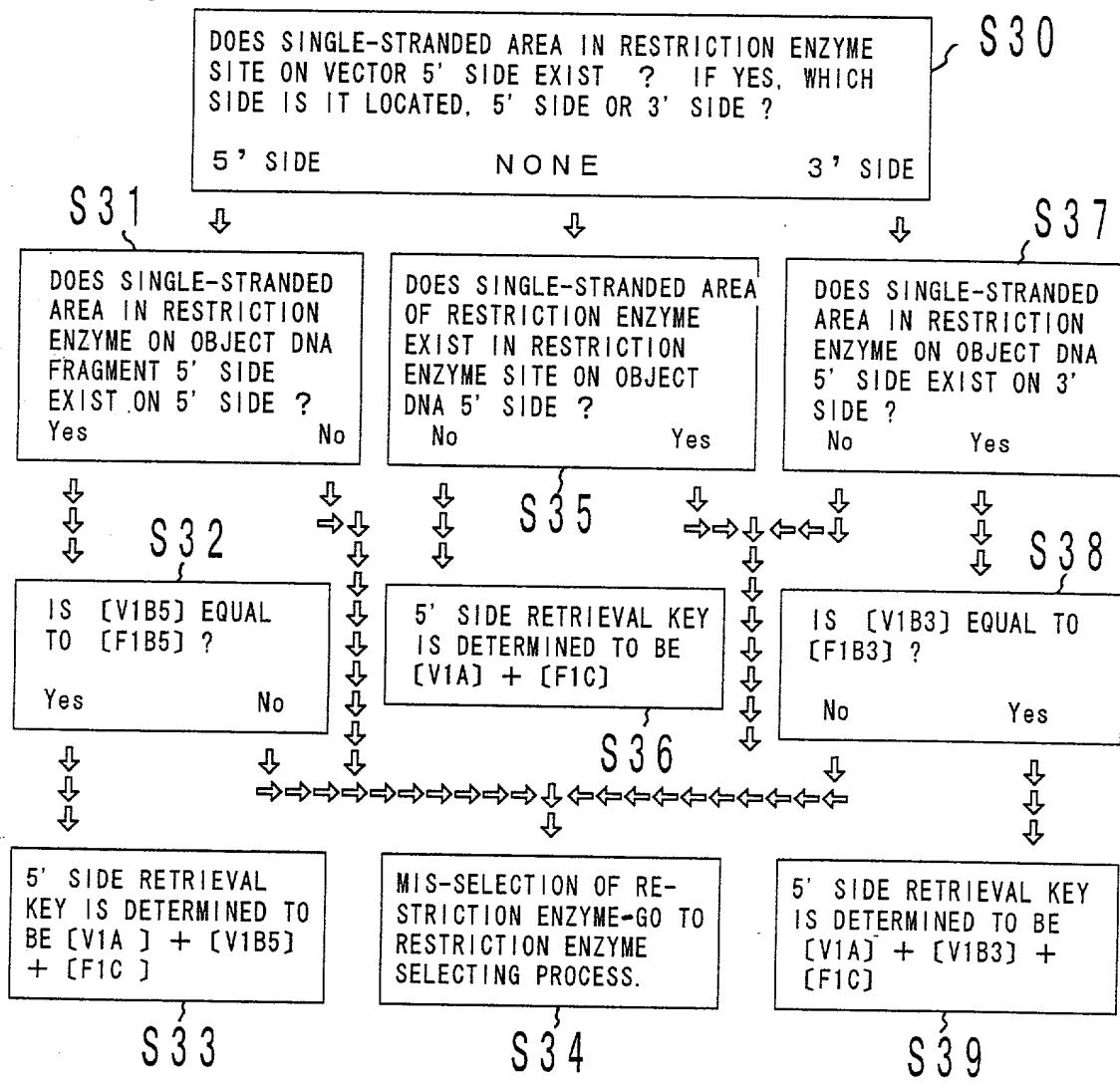
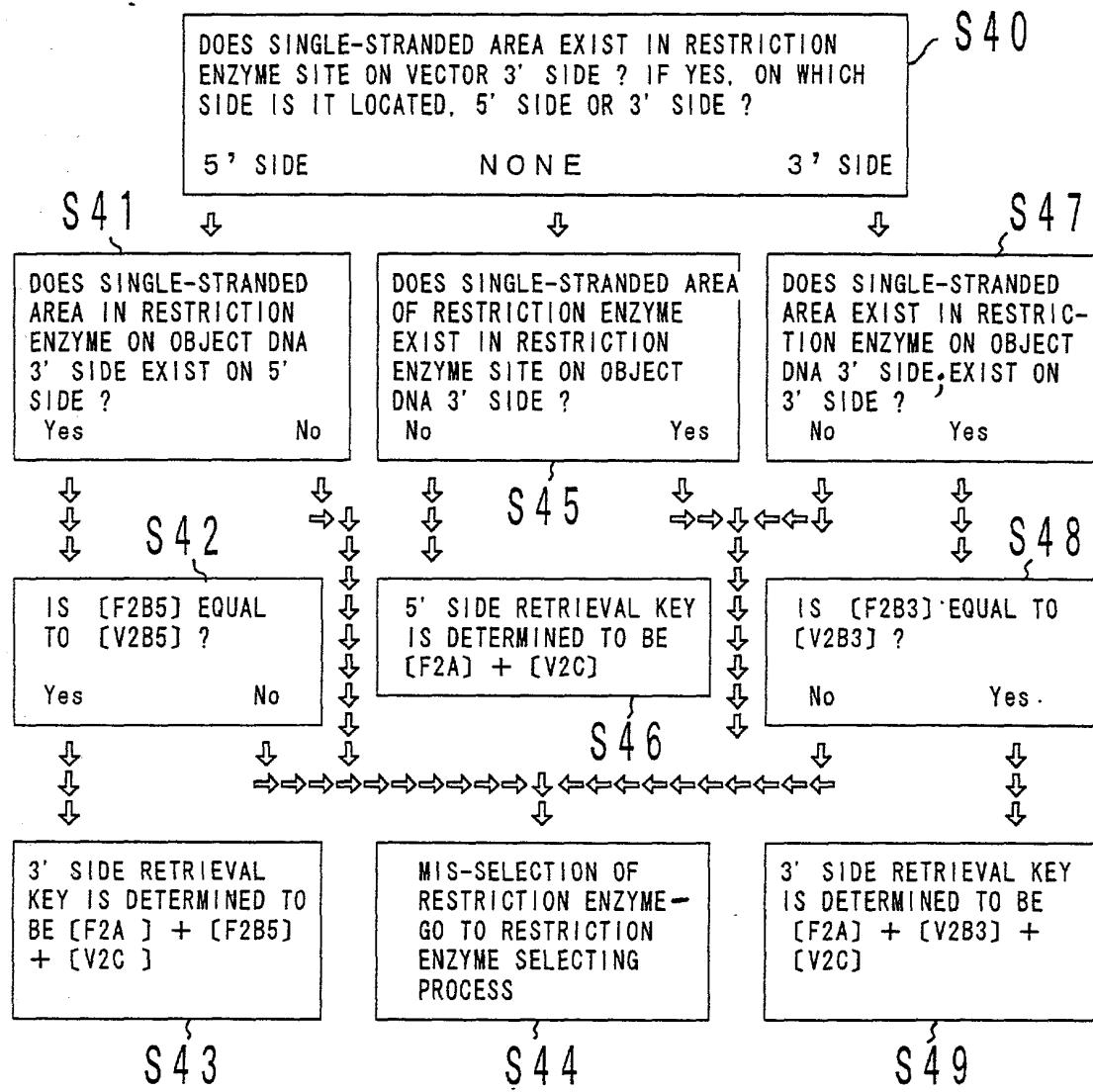


FIG. 15



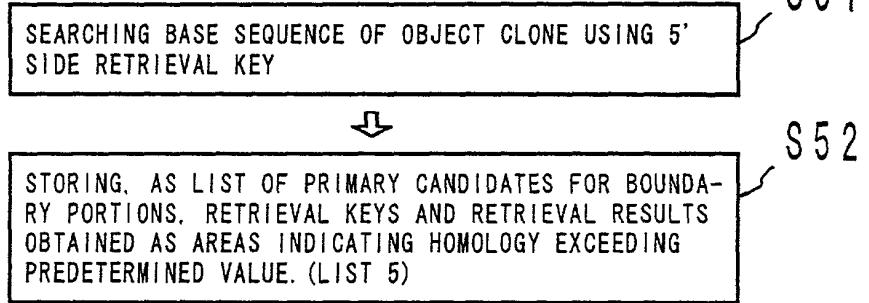
F I G. 16

WHEN HIND III IS SPECIFIED ON VECTOR 5' SIDE
XBA I IS SPECIFIED ON VECTOR 3' SIDE, HIND III IS
SPECIFIED ON OBJECT DNA 5' SIDE , AND XBA I IS
SPECIFIED ON OBJECT DNA 3' SIDE

(***** INDICATES RESIDUAL MULTIPLECLONING SITE
(+++++ INDICATES AN OBJECT DNA FRAGMENT

GTGCCAAGCTT++++++TCTAGAGGATCCCCGGGTACCGAGCTCGAATTCTGAAT
AAGCTT ↑
 ↑
5' SIDE RETRIEVAL KEY 9' SIDE RETRIEVAL KEY
(IN THIS EXAMPLE, (IN THIS EXAMPLE, XBA I SITE)
HIND III SITE)

FIG. 17



F I G. 18

SEARCHING BASE SEQUENCE OF OBJECT CLONE USING 3'
SIDE RETRIEVAL KEY

S 54

STORING, AS LIST OF PRIMARY CANDIDATES FOR BOUNDARY PORTIONS, RETRIEVAL KEYS AND RETRIEVAL RESULTS OBTAINED AS AREAS INDICATING HOMOLOGY EXCEEDING PREDETERMINED VALUE. (LIST 3)

S 55

F I G. 19

DEFINING, IN MULTIPLE CLONING SITE OF VECTOR, RESTRICTION ENZYME SITE USED IN SHEARING 5' SIDE IN MULTIPLE CLONING SITE OF VECTOR AND AREA OUTSIDE ON 5' SIDE AS 5' SIDE RESIDUAL MULTIPLE CLONING SITE (5MCS)

S 6 1

WHEN VECTOR DB CONTAINS BASE SEQUENCE OTHER THAN MULTIPLE CLONING SITE, SUM OF 5MCS AND 5 BASES ON 5' SIDE FROM 5MCS IS DEFINED AS 5' SIDE RESIDUAL VECTOR AREA (5VA). IF VECTOR DB CONTAINS ONLY BASE SEQUENCE OF MULTIPLE CLONING SITE IN VECTOR DB, THEN 5MCS IS 5VA.

S 6 2

A HOMOLOGY CHECK IS MADE ACCORDING TO FOLLOWING FLOWCHART
[ON ALL ELEMENTS IN PRIMARY CANDIDATES FOR BOUNDARY PORTIONS
(LIST 5) OBTAINED IN 5' SIDE HOMOLOGY RETRIEVAL]

DEFINING EACH CANDIDATE IN LIST 5 AND SEQUENCE AREA OUTSIDE ON 5' SIDE AS HOMOLOGY CHECK AREA (5HCA) FOR CORRESPONDING CANDIDATE

S 6 3

COMPARING NUMBER OF BASES IN 5' SIDE RESIDUAL VECTOR AREA (5VA), NUMBER OF BASES OF 5HCA, AND NUMBER OF BASES 20, AND DEFINING SMALLEST NUMBER OF BASES AS NUMBER OF BASES FOR USE IN HOMOLOGY CHECK (HCB)

S 6 4

EXTRACTING HCB BASES FROM 3' SIDE OF 5VA TO CHECK HOMOLOGY TO HCB BASES ON 3' SIDE OF 5HCA

S 6 5

WHEN CONSTANT HOMOLOGY IS OBTAINED, EXTRACTED BASES ARE DEFINED AS SECONDARY CANDIDATES FOR 5' SIDE BOUNDARY PORTIONS.

S 6 6

F I G . 2 0

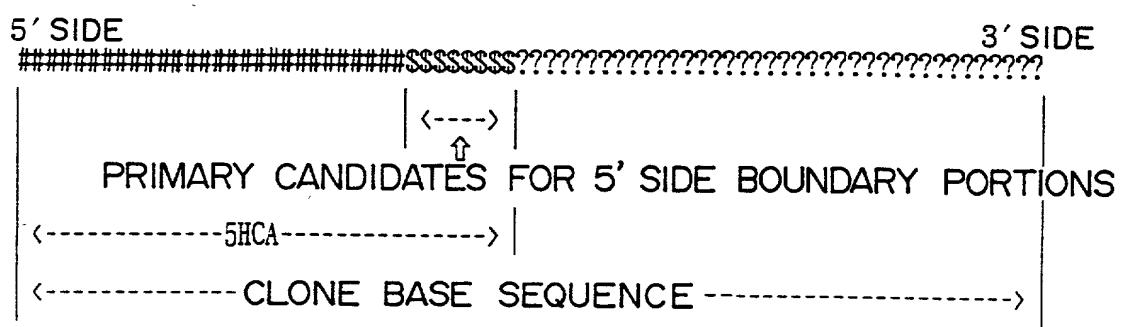
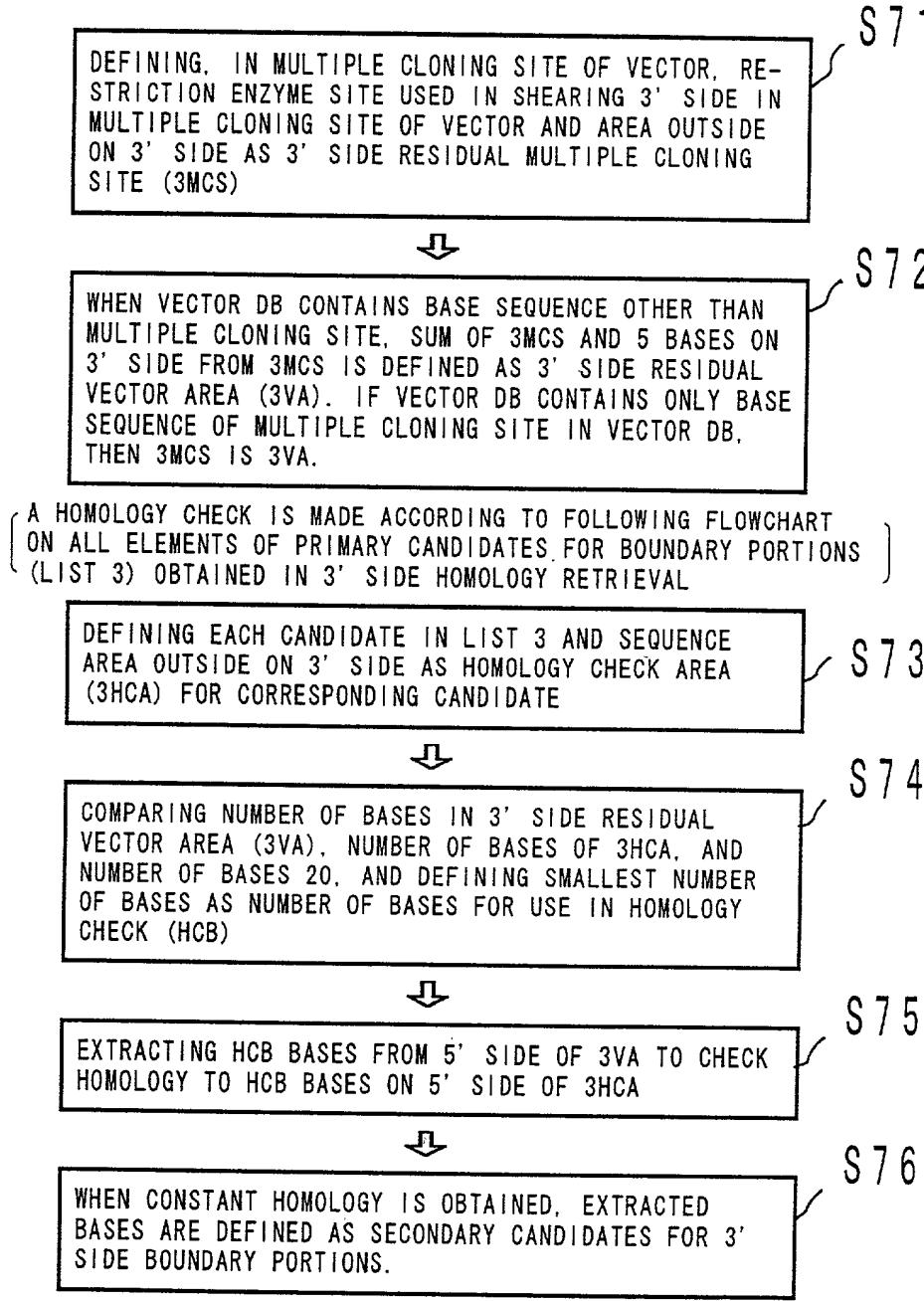


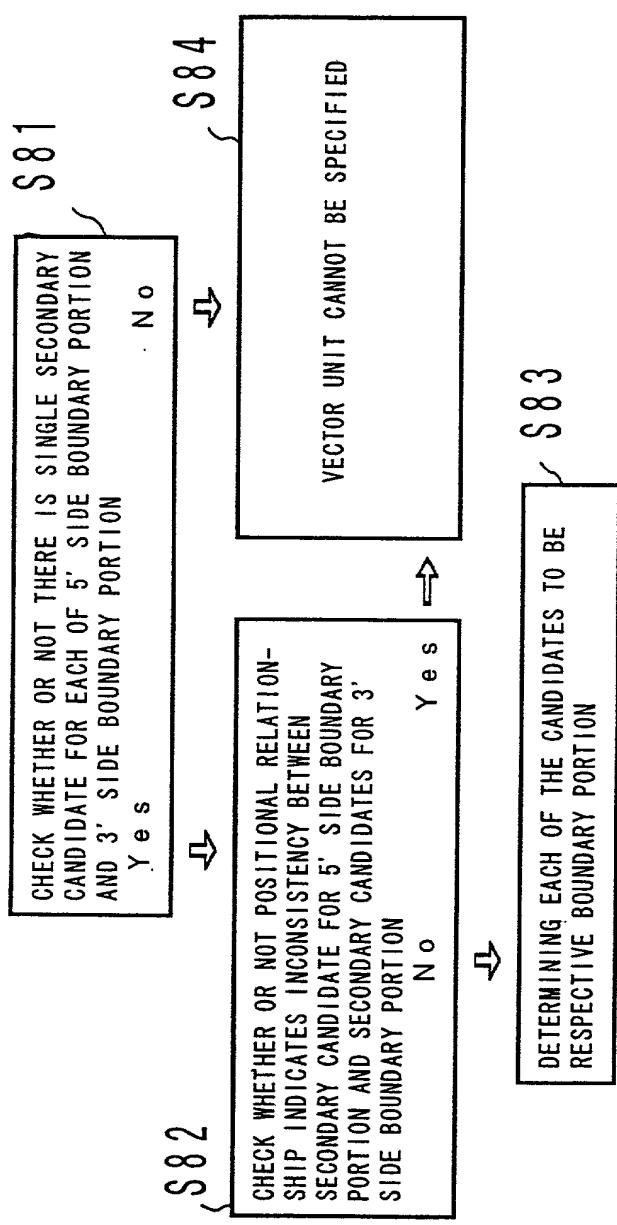
FIG. 21

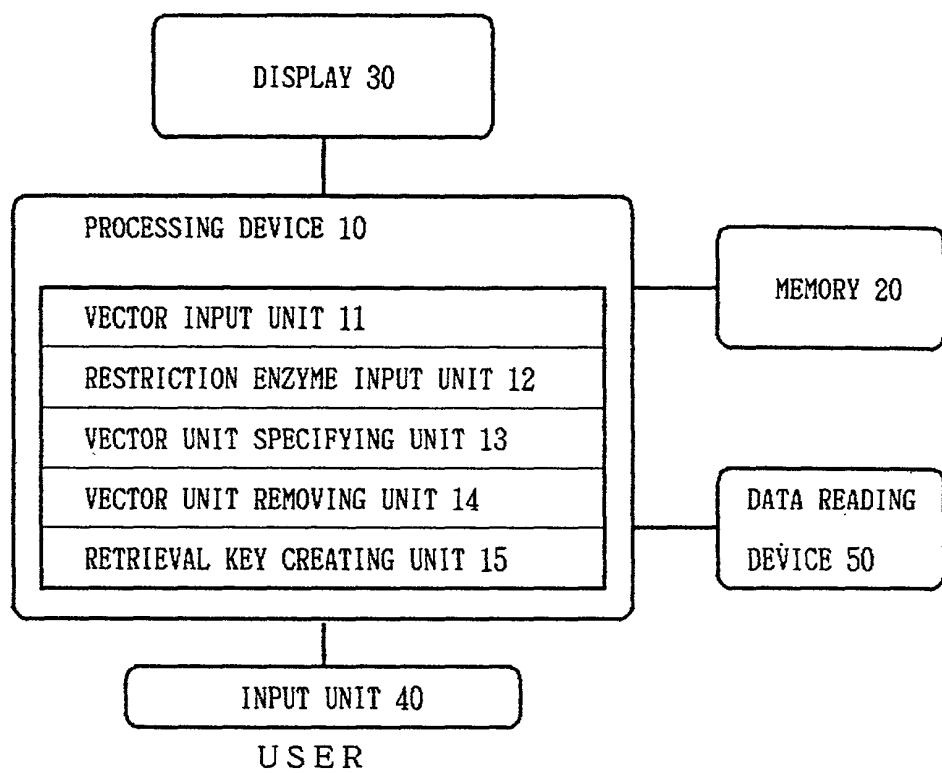


F I G. 22

FIG. 23

F I G. 24





F I G. 2 5